

## Skeletal muscle IL-15/IL-15R $\alpha$ and myofibrillar protein synthesis after resistance exercise

Pérez López, Alberto; McKendry, James; Martin-Rincon, Marcos ; Morales-Alamo, David; Pérez-Köhler, Bárbara ; Valadés, David; Buján, Julia ; Calbet, Jose; Breen, Leigh

DOI:  
[10.1111/sms.12901](https://doi.org/10.1111/sms.12901)

License:  
None: All rights reserved

Document Version  
Peer reviewed version

Citation for published version (Harvard):  
Pérez López, A, McKendry, J, Martin-Rincon, M, Morales-Alamo, D, Pérez-Köhler, B, Valadés, D, Buján, J, Calbet, J & Breen, L 2017, 'Skeletal muscle IL-15/IL-15R $\alpha$  and myofibrillar protein synthesis after resistance exercise', *Scandinavian Journal of Medicine and Science in Sports*. <https://doi.org/10.1111/sms.12901>

[Link to publication on Research at Birmingham portal](#)

### Publisher Rights Statement:

Checked for eligibility: 27/04/2017  
"This is the peer reviewed version of the following article: Pérez-López A, McKendry J, Martin-Rincon M, et al. Skeletal muscle IL-15/IL-15R $\alpha$  and myofibrillar protein synthesis after resistance exercise. Scand J Med Sci Sports. 2017, which has been published in final form at <https://doi.org/10.1111/sms.12901>. This article may be used for non-commercial purposes in accordance with Wiley Terms and Conditions for Self-Archiving."

### General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

### Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact [UBIRA@lists.bham.ac.uk](mailto:UBIRA@lists.bham.ac.uk) providing details and we will remove access to the work immediately and investigate.



### **Skeletal muscle IL-15/IL-15Ra and myofibrillar protein synthesis after resistance exercise**

Journal:	<i>Scandinavian Journal of Medicine and Science in Sports</i>
Manuscript ID	SJMSS-O-694-16.R2
Manuscript Type:	Original Article
Date Submitted by the Author:	27-Mar-2017
Complete List of Authors:	<p>Pérez-López, Alberto; University of Alcalá, Medicine and Medical Specialties. Faculty of Medicine and Health Sciences. Biomedical Research Networking Centre on Bioengineering, Biomaterials and Nanomedicine (CIBER-BBN); University of Alcalá, Biomedical Sciences. Faculty of Medicine and Health Sciences; University of Birmingham, School of Sport, Exercise and Rehabilitation Sciences; University of Las Palmas de Gran Canaria, Department of Physical Education. Faculty of Physical Education</p> <p>McKendry, James; University of Birmingham, School of Sport, Exercise and Rehabilitation Sciences; University of Birmingham, MRC-ARUK Centre for Musculoskeletal Ageing Research</p> <p>Martin-Rincon, Marcos; University of Las Palmas de Gran Canaria, Department of Physical Education. Faculty of Physical Education; University of Las Palmas de Gran Canaria, Research Institute of Biomedical and Health Sciences (IUIBS)</p> <p>Morales-Alamo, David; University of Las Palmas de Gran Canaria, Department of Physical Education. Faculty of Physical Education; University of Las Palmas de Gran Canaria, Research Institute of Biomedical and Health Sciences (IUIBS)</p> <p>Pérez-Köhler, Bárbara; University of Alcalá, Department of Medicine and Medical Specialties. Faculty of Medicine and Health Sciences. Biomedical Research Networking Centre on Bioengineering, Biomaterials and Nanomedicine (CIBER-BBN)</p> <p>Valadés, David; University of Alcalá, Department of Biomedical Sciences. Faculty of Medicine and Health Sciences.</p> <p>Buján, Julia; University of Alcalá, Department of Medicine and Medical Specialties. Faculty of Medicine and Health Sciences. Biomedical Research Networking Centre on Bioengineering, Biomaterials and Nanomedicine (CIBER-BBN)</p> <p>Calbet, Jose; University of Las Palmas de Gran Canaria, Department of Physical Education. Faculty of Physical Education; University of Las Palmas de Gran Canaria, Research Institute of Biomedical and Health Sciences (IUIBS)</p> <p>Breen, Leigh; University of Birmingham, School of Sport, Exercise and Rehabilitation Sciences; University of Birmingham, MRC-ARUK Centre for Musculoskeletal Ageing Research</p>
Keywords:	Myokines, IL-15/IL-15Ra axis, strength training, muscle protein synthesis/breakdown

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60



SCHOLARONE™  
Manuscripts

PROOF

# **Skeletal muscle IL-15/IL-15R $\alpha$ and myofibrillar protein synthesis after resistance exercise**

Alberto Pérez-López<sup>1,2,3,4</sup>, James McKendry<sup>3,5</sup>, Marcos Martin-Rincon<sup>4,6</sup>, David Morales-Alamo<sup>4,6</sup>, Bárbara Pérez-Köhler<sup>1</sup>, David Valadés<sup>2</sup>, Julia Buján<sup>1</sup>, Jose A. L. Calbet<sup>4,6</sup> and Leigh Breen<sup>3,5</sup>.

<sup>1</sup>Department of Medicine and Medical Specialities. Faculty of Medicine and Health Sciences. University of Alcalá, Madrid, Spain. Biomedical Research Networking Centre on Bioengineering, Biomaterials and Nanomedicine (CIBER-BBN), Madrid, Spain.

<sup>2</sup>Department of Biomedical Sciences. Faculty of Medicine and Health Sciences. University of Alcalá, Madrid, Spain.

<sup>3</sup>School of Sport, Exercise and Rehabilitation Sciences. University of Birmingham, Birmingham, UK.

<sup>4</sup>Department of Physical Education. Faculty of Physical Education. University of Las Palmas de Gran Canaria, Las Palmas de Gran Canaria, Spain.

<sup>5</sup>MRC-ARUK Centre for Musculoskeletal Ageing Research. University of Birmingham, Birmingham, UK.

<sup>6</sup>Research Institute of Biomedical and Health Sciences (IUIBS). Las Palmas de Gran Canaria, Spain.

## **Address for correspondence:**

Alberto Pérez-López

Department of Medicine and Medical Specialties.

Faculty of Medicine and Health Sciences, University of Alcalá.

Ctra. Madrid-Barcelona km 33,600

28871 Alcalá de Henares, Madrid (Spain)

E-mail: [Alberto\\_perez-lopez@hotmail.com](mailto:Alberto_perez-lopez@hotmail.com)

Phone: (+34) 636 710 130

**Running head:** IL-15/IL-15R $\alpha$  in Resistance Exercise

**ABSTRACT**

*In vitro* and *in vivo* studies described the myokine IL-15 and its receptor IL-15R $\alpha$  as anabolic/anti-atrophy agents, however the protein expression of IL-15R $\alpha$  has not been measured in human skeletal muscle and data regarding IL-15 expression remain inconclusive. The purpose of the study was to determine serum and skeletal muscle IL-15 and IL-15R $\alpha$  responses to resistance exercise session and to analyse their association with myofibrillar protein synthesis (MPS). Fourteen participants performed a bilateral leg resistance exercise composed of 4 sets of leg press and 4 sets of knee extension at 75% 1RM to task failure. Muscle biopsies were obtained at rest, 0, 4 and 24h post-exercise and blood samples at rest, mid-exercise, 0, 0.3, 1, 2, 4 and 24h post-exercise. Serum IL-15 was increased by ~5.3-fold immediately post-exercise, while serum IL-15R $\alpha$  decreased ~75% over 1h post-exercise ( $P<0.001$ ). Skeletal muscle IL-15R $\alpha$  mRNA and protein expression were increased at 4h post-exercise by ~2-fold ( $P<0.001$ ) and ~1.3-fold above rest ( $P=0.020$ ), respectively. At 24h post-exercise IL-15 ( $P=0.003$ ) and IL-15R $\alpha$  mRNAs increased by ~2-fold ( $P=0.002$ ). Myofibrillar fractional synthetic rate between 0-4h was associated with IL-15R $\alpha$  mRNA at rest ( $r=0.662$ ,  $P=0.019$ ), 4h ( $r=0.612$ ,  $P=0.029$ ) and 24h post-exercise ( $r=0.627$ ,  $P=0.029$ ). Finally, the muscle IL-15R $\alpha$  protein up-regulation was related to Leg press 1RM ( $r=0.688$ ,  $P=0.003$ ) and total weight lifted ( $r=0.628$ ,  $P=0.009$ ). In conclusion, IL-15/IL-15R $\alpha$  signalling pathway is activated in skeletal muscle in response to a session of resistance exercise.

**Keywords:** Myokines, IL-15/IL-15R $\alpha$  axis, strength training, muscle protein synthesis/breakdown.

## 53 INTRODUCTION

54 Interleukin-15 (IL-15) and its cognate receptor alpha (IL-15R $\alpha$ ) have been implicated in the  
55 regulation of anabolic/catabolic balance of human skeletal muscle (Busquets et al., 2005;  
56 Furmanczyk & Quinn, 2003; Pistilli et al., 2007; Quinn et al., 2002; Quinn et al., 1995;  
57 Riechman et al., 2004). However, most of the evidence is indirect and the protein expression of  
58 IL-15R $\alpha$  has not been determined in human skeletal muscle.

59 IL-15 is a pleiotropic cytokine member of the 4 alpha-helix bundle family (Grabstein et  
60 al., 1994). IL-15 has been shown to stimulate protein accretion and myosin heavy chain (MHC)  
61 accumulation in differentiated myocytes (Quinn et al., 1995) and myotubes (Furmanczyk &  
62 Quinn, 2003; Quinn et al., 2002), while reducing protein degradation (Quinn et al., 2002). In  
63 humans, circulating IL-15 is elevated in response to a single session of resistance exercise in  
64 untrained and trained states (Riechman et al., 2004). In agreement with a muscular origin, IL-15  
65 mRNA was increased 2-fold in *vastus lateralis* muscle 24h after a bilateral leg press and knee  
66 extension resistance exercise session, although this was not accompanied by a change in  
67 circulating or muscular IL-15 protein expression (Nielsen et al., 2007). Therefore, despite the  
68 fact that *in vitro* studies indicate a role for skeletal muscle IL-15 in anabolism, studies in  
69 humans are inconclusive.

70 Although part of the effects of IL-15 are mediated by its binding to IL-15R $\alpha$  (Dubois et  
71 al., 2002; Duitman et al., 2008; Sato et al., 2007), this alpha-receptor may also exert functions  
72 independent from IL-15 in skeletal muscle. IL-15R $\alpha$  may have a role in determining the  
73 phenotype and fatigability of muscle fibers, and mitochondrial fuel utilization (Loro et al.,  
74 2015; O'Connell et al., 2015; O'Connell et al., 2015). In addition, human studies indicate that  
75 IL-15R $\alpha$  may be involved in muscle hypertrophy and strength gains after resistance training  
76 (Pistilli et al., 2008; Riechman et al., 2004). In this regard, two single nucleotide polymorphism  
77 (SNPs) in exon 7 and 4 of the IL-15R $\alpha$  could explain part of the variability in the hypertrophy  
78 observed after 10 weeks of whole-body resistance training (Riechman et al., 2004), whereas IL-  
79 15R $\alpha$  SNPs, rs2296135 and rs22228059, were positively associated with pre- and post-exercise

isometric strength and muscle volume, respectively, after 12 weeks of resistance training of the flexor-extensor muscles of the elbow (Pistilli et al., 2008).

Despite the potential implication of IL-15 and IL-15R $\alpha$  in skeletal muscle anabolic/catabolic balance, direct evidence is lacking as no human study has determined whether changes in skeletal muscle IL-15R $\alpha$  mRNA and protein expression are associated with protein synthesis. Recently, we reported a ~2-fold elevated myofibrillar protein synthesis (MPS) response during the first four hours (0-4h) after a single session of resistance exercise in healthy young males (McKendry et al., 2016), however, skeletal muscle IL-15 and IL-15R $\alpha$  were not determined. Therefore, the aim of this study was to determine whether circulating and skeletal muscle IL-15 and IL-15R $\alpha$  might have a role in the regulation of myofibrillar protein synthesis, after a single session of resistance exercise.

We hypothesised that skeletal muscle IL-15 and IL-15R $\alpha$  expressions would be up-regulated after a single session of resistance exercise and that IL-15 and IL-15R $\alpha$  expression in skeletal muscle would be associated with myofibrillar protein synthesis.

**MATERIALS AND METHODS**

**Participants**

A full description of the methods, study design and participant characteristics, from which part of current data are drawn, has been previously published (McKendry et al., 2016). Volunteers were aged from 18 to 35 years and had been participating in resistance training programmes > 2 days/week during  $\geq$  1 year prior to start of current study. Subjects' characteristics are presented in Table 1. Prior to study enrolment all procedures were explained to participants who then gave their written informed consent. Ethical approval was obtained through the NHS Black Country Research Ethics Committee (13/WM/0455) in accordance with the latest version (7<sup>th</sup>) of the Declaration of Helsinki.

## 106 **Experimental design**

107 All participants reported to the School of Sport, Exercise and Rehabilitation Sciences  
108 (SportExR) laboratory on 3 separate occasions. During visit one, participants underwent  
109 preliminary assessments of body composition and maximal leg strength. Then, within a period  
110 of 8 days, the volunteers returned to the laboratory for the experimental trial, which consisted of  
111 a single session of bilateral lower-limb resistance exercise with muscle biopsies obtained at  
112 baseline, immediately after (0h), 4 and 24h post-exercise, and blood samples at baseline, mid-  
113 exercise, and 0, 0.3, 1, 2, 4 and 24h post-exercise to assess the systemic and skeletal muscle  
114 responses of the IL-15/IL-15R $\alpha$  axis. As part of the original investigation (McKendry et al.,  
115 2016), participants were matched in pairs based on anthropometric, strength and training  
116 characteristics before to be randomly allocated to either 1-min or 5-min of passive rest between  
117 resistance exercise sets. Since no significant differences were observed in circulating or  
118 intramuscular IL-15 and IL-15R $\alpha$  measurements between the 1-min (N = 7) and 5-min groups  
119 (N = 7), participants were treated as a single group for the purpose of the present analyses. Of  
120 the 16 participants included in the original study, 14 were analysed in the present investigation  
121 due to insufficient muscle tissue in the two subjects excluded.

122

## 123 **Experimental protocol**

124 A detailed description of the experimental protocol and analytical methods can be found  
125 elsewhere (McKendry et al., 2016). Briefly, regional and whole-body composition was  
126 determined by dual energy x-ray absorptiometry (Discovery DXA Systems, Hologic Inc.,  
127 Bedford, MA, USA). Thereafter, one-repetition maximum (1RM) strength during leg press and  
128 knee extension was assessed (Cybex VR-3, MA, USA). Approximately seven days later  
129 participants reported to the laboratory at 07.00 hours being fasted for 10-12h. Upon arrival, a  
130 cannula was inserted into a forearm vein to obtain arterialised blood samples into a tube  
131 prepared for serum separation (BD, Oxford, UK). After resting supine in bed for 2.5h, a muscle  
132 biopsy was obtained from the *vastus lateralis* of one leg (~120mg of tissue) (Bergstrom, 1975),



under local anaesthesia (1% lidocaine). Skeletal muscle sample was cleaned from any fat or connective tissue before being frozen in liquid nitrogen. Following the muscle biopsy, participants completed a session of bilateral lower-extremity resistance exercise on leg press and knee extension machines. Exercise consisted of four sets of 8-15 repetitions per exercise at 75% of 1RM, each set performed to task failure. At the end of the last repetition, a second muscle biopsy was obtained ~3 cm proximal from the first biopsy through a new incision. Immediately after the second biopsy, the volunteers ingested 25 g of whey protein isolate (MyProtein, Cheshire, UK) dissolved in 400 mL of water. During the next four hours, participants rested supine and then a third muscle biopsy was obtained from a new incision, ~3 cm proximal to the second biopsy. The following morning at 7.00 h, participants returned to the laboratory after a 10-12h overnight fast and a cannula was inserted into a forearm vein to obtain a blood sample followed by the fourth and last biopsy, which was obtained from the *vastus lateralis* of the contralateral leg.

The participants received three standardised meals for consumption the evening prior to the experimental trial, as well as the afternoon and evening after the experimental trial. The diet was composed by ~97 g of CHO (~58%), ~34 g of protein (~20%) and ~37 g of fat (~22%) with an energy content of ~871 kcal per meal. Consumption of ethanol or caffeine was not allowed 24h before the experiments neither during the study.

**Blood Analysis**

***Serum IL-15 and IL-15Rα***

After collection, all blood samples were centrifuged for 15 minutes at 1000 g, aliquoted and stored at -80 °C. Two high-sensitivity enzyme-linked immunosorbent assay (ELISA) kits were used to determine the serum concentration of IL-15 and IL-15Rα in duplicates. IL-15 was measured using Human IL-15 Quantikine ELISA kit (R&D Systems, MN, USA) recognizing both natural and recombinant human IL-15 (range: 0.49 – 62.5 pg/mL; intra- and inter-assay

coefficients of variation (CV) were 4.2 and 7.4%, respectively). Serum IL-15R $\alpha$  was measured using Human IL-15 receptor subunit alpha ELISA kit (Wuhan EIAab Science, Wuhan, China) recognizing both natural and recombinant human IL-15R $\alpha$  (range: 0.49 – 62.5 pg/mL; intra- and inter-assay CVs were 4.4 and 7.8%, respectively).

## Muscle Tissue Analysis

### *Protein extraction and Western Blot procedures.*

Approximately 25-30 mg of muscle tissue was powdered on dry ice using a Cellcrusher<sup>TM</sup> tissue pulveriser (Cellcrusher limited, Cork, Ireland) and a sucrose lysis buffer was used to prepare the samples for Western Blot as previously described (Philp et al., 2011). Equal amounts of protein (35  $\mu$ g per sample) were boiled for 5 min in 1 x Laemmli sample buffer, separated on 10% SDS-PAGE gels (Bio-Rad, Copenhagen, Denmark) for 45 min and transferred to polyvinylidene difluoride (PVDF) membranes at constant voltage and 0.4 A for 1.5 h. Subsequently, membranes were incubated overnight with primary antibodies against IL-15 (sc-7889) and IL-15R $\alpha$  (sc-271366), purchased from Santa Cruz Biotechnology (Dallas, USA). Both antibodies were diluted into BSA-blocking buffer containing 4% bovine serum albumin in Tris-buffered saline with 0.1% Tween 20. Antibody specific labelling was revealed by incubation with an HRP-conjugated goat anti-rabbit (IL-15) or anti-mouse (IL-15R $\alpha$ ) antibodies (1:5000), both diluted in 5% blotto blocking buffer and visualised with ECL Western blotting detection system using a ChemiDoc XRS (Bio-Rad, Copenhagen, Denmark). Imaging and band quantification were performed using the Quantity One 1-D Analysis software (Bio-Rad, Copenhagen, Denmark). Test samples were run together with a control sample from a subject who did not take part in the study. The control sample was loaded in three different lanes and used as an internal control for inter-gel variability. Overall, the mean CVs of the controls were 13.9% (IL-15) and 12.3% (IL-15R $\alpha$ ). Control samples and a total protein staining-technique method (reactive brown) were used to accurate protein quantification for loading control.

***RNA Isolation and quantitative real-time reverse transcription polymerase chain reaction (qRT-PCR).***

Approximately 15-20 mg of skeletal muscle tissue was used for the RNA isolation. The RNA was extracted by guanine-phenol-chloroform isothiocyanate procedures using TRIzol (Invitrogen, Carlsbad, CA, USA). Then, RNA was recovered from the aqueous phase by precipitation; the amount and purity was measured by optical density at 260/280 nm and 260/230 nm in a NanoDrop ND-100 spectrophotometer (Thermo Fisher Scientific Inc., DE, USA).

Reverse transcription was performed to synthesize cDNA from 200 ng of the total RNA using Oligo dT primers (GE Healthcare Bio-Sicences, Buckinghamshire, UK) and M-MLV reverse transcriptase enzyme (Invitrogen, Carlsbad, CA, USA). The cDNA was amplified using the primers presented in Table 2 (self-designed and tested in skeletal muscle from human donors, data not shown). The qRT-PCR mixture was composed by 5 µL of the inverse transcription product (cDNA) diluted 1:20, 10 µL of iQ SYBR Green Supermix (Bio-Rad, Copenhagen, Denmark) and 1 µL (6 mM) of the primer selected. The final reaction volume (20 µL) was used to perform the qRT-PCR in a StepOnePlus Real-Time PCR System (Applied Biosystems, Foster City, CA, USA). All samples were subjected to an initial stage of 10 min at 95°C. The conditions for PCR amplification were as follows: 45 cycles of 95 °C for 15 s, 60 °C for 30 s and 72 °C for 1 minute, for both IL-15 and IL-15Rα. Finally, mRNA expressions of IL-15 and IL-15Rα were determined in triplicates, and normalized using β-actin and glyceraldehyde 3-phosphate-dehydrogenase (GAPDH) as housekeeping genes. β-actin and GAPDH expression remained unchanged.

***Myofibrillar and plasma tracer enrichment.***

Procedures for muscle myofibrillar protein isolation, plasma-free amino acid extraction and <sup>13</sup>C<sub>6</sub> phenylalanine enrichment, and calculation of myofibrillar fractional synthetic rate (FSR) at rest, 0-4h and 24-28h post-exercise are described in McKendry et al. (2016).

Briefly, a primed continuous infusion of L-[ring- $^{13}\text{C}_6$ ]phenylalanine (prime,  $2\ \mu\text{mol kg}^{-1}$ ; infusion,  $0.05\ \mu\text{mol kg}^{-1}\ \text{min}^{-1}$ ; Cambridge Isotope Laboratories, Andover, MA, USA) was implemented during both experimental trial days in conjunction with muscle biopsy and blood sampling. In both experimental days, the infusion was initiated immediately after the drawn of the first blood sample ( $\sim 7.05\ \text{h}$ ) and finished when the last muscle biopsy sample of the day was obtained ( $7.5\ \text{h}$  and  $5.5\ \text{h}$  after the beginning of the infusion on day 1 and 2, respectively). Upon thawing, plasma samples were purified on cation-exchange columns. The amino acids were then converted to their N-tert-butyldimethylsilyl-N-methyltrifluoroacetamide (MTBSTFA) derivative. Plasma [ $^{13}\text{C}_6$ ]phenylalanine enrichment was determined by gas chromatography–mass spectrometry (GCMS; model 5973; Hewlett Packard, Palo Alto, CA, USA) by monitoring ions 234/240. The myofibrillar protein fraction was extracted and hydrolysed overnight. Constituent amino acids in the myofibrillar fraction were purified on cation-exchange columns. Amino acids in the myofibrillar fraction were then converted to their N-acetyl-n-propyl ester derivative. Plasma [ $^{13}\text{C}_6$ ]phenylalanine enrichment was determined by gas chromatography–mass spectrometry (GC-C-IRMS; Delta-plus XP; Thermofinnigan, Hemel Hempstead, UK) by monitoring ions 44/45. Pre-infusion and mean plasma [ $^{13}\text{C}_6$ ]phenylalanine enrichment were used as a proxy for basal muscle protein enrichment and to determine an “estimated” intracellular precursor enrichment, respectively. The fractional synthesis rate (FSR) of the myofibrillar protein fraction was calculated from the incorporation of [ $^{13}\text{C}_6$ ]phenylalanine into protein using the standard precursor-product model (Wolfe & Chinkes, 2005).

### Statistical Analysis

Data collected in the study were analysed using the statistical package SPSS v. 22.0 (SPSS Inc., Chicago, IL, USA), and Graph Prism 6 (GraphPad software, Inc. La Jolla, CA, USA). Firstly, a Shapiro-Wilks was used to test the normality of the data ( $P > 0.05$ ). Subsequently, non-normally distributed variables were logarithmically transformed. Circulating and skeletal muscle expression of IL-15 and IL-15R $\alpha$  were analysed between groups (1- vs. 5-min rest)

241 using a two-way, repeated measures ANOVA (time x condition). The area under the curve  
242 (AUC) was determined using trapezoid method and compared between groups using a paired  
243 Student's t-test. Since no significant differences were observed between the 1- and 5-min  
244 recovery, both groups were combined for further analyses.

245 To determine time effects of the intervention on serum, protein and mRNA levels of IL-  
246 15 and IL-15R $\alpha$ , ANOVA for repeated measures was performed. Tukey HSD correction was  
247 used as *post-hoc* test when significant differences were detected. Finally, linear regression  
248 analysis was carried out to test the potential associations between skeletal muscle and  
249 circulating levels of IL-15 and IL-15R $\alpha$ , as well as between the former and resistance training  
250 variables, body composition and myofibrillar FSR. The effect of size (ES) was calculated as eta  
251 squared statistic ( $\eta^2$ ) to verify time, condition and between groups differences in systemic and  
252 intramuscular IL-15 and IL-15R $\alpha$  expression. Values are reported as mean  $\pm$  standard deviation  
253 (SD); a  $P < 0.05$  was considered statistically significant.

255 **RESULTS**

256 Variables describing the resistance exercise session performed by the subjects are presented in  
257 Table 3.

259 **IL-15 response to a single dose of resistance exercise.**

260 Skeletal muscle mRNA and protein expression levels, and serum IL-15 concentrations in  
261 response to a single session of resistance exercise are illustrated in Figure 1. A progressive  
262 increase in mRNA expression was found following resistance exercise ( $P = 0.002$ ,  $ES = 0.35$ ),  
263 reaching statistical significance at 4h ( $P = 0.019$ ,  $ES = 0.37$ ) and 24h post-exercise, where a 2-  
264 fold elevation above pre-exercise resting values was found ( $P = 0.003$ ,  $ES = 0.44$ ; Fig. 1A). No  
265 significant changes were observed in IL-15 muscle protein expression above pre-exercise  
266 resting values ( $P = 0.563$ ,  $ES = 0.15$ ; Fig. 1B). Serum IL-15 concentration increased  
267 significantly above pre-exercise resting values during the post-exercise period ( $P = 0.001$ ,  $ES =$

0.46; Fig. 1C), peaking immediately post-exercise ( $P < 0.001$ ,  $ES = 0.31$ ), and remaining elevated at 24h post-exercise ( $P = 0.001$ ,  $ES = 0.42$ ).

### **IL-15R $\alpha$ response to a single dose of resistance exercise.**

Skeletal muscle mRNA and protein expression levels, and serum IL-15R $\alpha$  concentration in response to a single session of resistance exercise are illustrated in Figure 2. A significant 2-fold increase in IL-15R $\alpha$  mRNA expression above pre-exercise resting values was observed at 4h ( $P < 0.001$ ,  $ES = 0.42$ ) and 24h post-exercise ( $P = 0.002$ ,  $ES = 0.35$ ; Fig. 2A). In contrast to IL-15, skeletal muscle protein expression of IL-15R $\alpha$  increased by 1.3-fold ( $P = 0.020$ ,  $ES = 0.34$ ; Fig. 2B) above pre-exercise resting values at 4h post-exercise, returning to baseline levels 24h post-exercise ( $P = 0.036$ ,  $ES = 0.32$ ). Despite of the lack of significant differences between inter-set rest period (1- vs. 5-min groups), the 5-min group tended to show an elevated protein and mRNA expression at 4h and post-exercise ( $\sim 10\%$ ;  $P = 0.103$ ,  $ES = 0.55$  and  $P = 0.092$ ,  $ES = 0.57$ , respectively; supplementary figure 2). Finally, compared to pre-exercise values, serum IL-15R $\alpha$  concentration was significantly reduced during the first 60 min following the training session ( $P < 0.001$ ;  $ES = 0.47$ ; Fig. 2C).

### **Correlation analysis**

#### ***IL-15/IL-15R $\alpha$ and resistance exercise variables.***

Leg Press 1RM strength was negatively associated with IL-15 serum concentration ( $r = -0.800$ ,  $P = 0.003$ ) but not with IL-15R $\alpha$  serum concentration. While the serum IL-15 concentration response to resistance exercise (AUC) was negatively associated with knee extension training volume ( $r = -0.637$ ,  $P = 0.042$ ) and time-under-tension (T-U-T) ( $r = -0.718$ ,  $P = 0.019$ ). Post-exercise (0h), serum IL-15 concentrations were associated with the total volume of knee extension exercise ( $r = -0.934$ ,  $P < 0.001$ ). Furthermore, skeletal muscle protein expression of IL-15R $\alpha$  at 4h post-exercise was associated positively with 1RM Leg press strength ( $r = 0.559$ ,  $P = 0.037$ ) and total training load ( $r = 0.628$ ,  $P = 0.009$ ).

295

296 *Skeletal muscle and circulating expressions of IL-15/IL-15Rα*

297 At baseline, serum IL-15 concentration was positively associated with intramuscular protein IL-

298 15 levels ( $r = 0.649$ ,  $P = 0.031$ ), but not with mRNA expression. Serum IL-15 immediately

299 post-exercise was associated with pre-exercise levels of skeletal muscle protein expression of

300 IL-15 and IL-15Rα ( $r = 0.582$ ,  $P = 0.037$ ;  $r = -0.599$ ,  $P = 0.031$ , respectively). At baseline, IL-

301 15 mRNA was associated with IL-15Rα mRNA ( $r = 0.592$ ,  $P = 0.043$ ), as well as immediately

302 ( $r = 0.791$ ,  $P = 0.005$ ) and 24h post-exercise ( $r = 0.653$ ,  $P = 0.021$ ). Additionally, IL-15 and IL-

303 15Rα mRNA expressions at 24h post-exercise was negatively associated with serum

304 concentration of IL-15 and IL-15Rα ( $r = -0.620$ ,  $P = 0.042$ ;  $r = -0.727$ ,  $P = 0.005$ ; respectively).

305

306 *Skeletal muscle IL-15/IL-15Rα expression and myofibrillar protein synthesis.*

307 The myofibrillar fractional synthetic rate (FSR) increased by ~2-fold above resting values from

308 0-4h post-exercise (McKendry et al., 2016) and was associated with IL-15Rα mRNA levels at

309 baseline ( $r = 0.662$ ,  $P = 0.019$ ), 4h ( $r = 0.612$ ,  $P = 0.029$ ) and 24h post-exercise ( $r = 0.627$ ,  $P =$

310  $0.029$ ) (Figure 3). Moreover, the delta changes ( $\Delta$ ), from pre-exercise to 4h post-exercise, of IL-

311 15Rα mRNA expression and FSR showed a tendency to be associated at 4h post-exercise ( $r =$

312  $0.481$ ;  $P = 0.096$ ). No association was observed between myofibrillar FSR and skeletal muscle

313 IL-15 mRNA or muscle protein expression of either IL-15 or IL-15Rα at any time.

314

315 **DISCUSSION**

316 The present study demonstrates that the gene and protein expression of IL-15Rα is up-regulated

317 in skeletal muscle after a single session of resistance training. The increase in myofibrillar

318 protein synthesis during 0-4h post-exercise was associated with the expression of IL-15Rα

319 mRNA at 4h, which occurred concomitantly with an increase of skeletal muscle IL-15Rα

320 protein levels, suggesting increased translation of the IL-15Rα gene. These findings indicate



that IL-15R $\alpha$  could have a role in mediating the increase in myofibrillar protein synthesis observed in skeletal muscle after a single session of resistance training.

Although, in our previous study we demonstrated that myofibrillar protein synthesis rates were greater when high volume, moderate-intensity resistance exercise was performed with long (5-min) compared with short (1-min) inter-set rest duration (McKendry et al., 2016), in the present study, we did not observe significant differences between groups in circulating or intramuscular IL-15 and IL-15R $\alpha$  expressions, despite skeletal muscle IL-15R $\alpha$  tended to be elevated in the 5-min compared to the 1-min group (supplementary figure 2). This lack of differences could be interpreted as evidence to refute the association between skeletal muscle IL-15/IL-15R $\alpha$  and MPS. Nevertheless, the effect sizes and statistical outputs ( $P < 0.10$ ) indicate that a potential difference between groups may actually exist. This suggestion is also supported by the fact that the association between IL-15R $\alpha$  and MPS in the early recovery phase (0-4h post-exercise) was observed in each group separately (1-min group,  $r = 0.592$ ,  $P = 0.052$ ; and 5-min,  $r = 0.684$ ,  $P = 0.043$ ). Therefore, our results provide the framework for future studies to further clarify whether the IL-15/IL-15R $\alpha$  response to strength training reported here have a physiologically relevant role in human skeletal muscle adaptation to this type of exercise.

The interleukin-15 subunit alpha-receptor (IL-15R $\alpha$ ) is a key subunit receptor of IL-15 that regulates its signalling in several cell types (Budagian et al., 2006; Dubois et al., 2002; Duitman et al., 2008; Sato et al., 2007). In addition to the common receptor-binding functions, IL-15R $\alpha$  has been shown to function by itself, without the need for IL-15 binding (Loro et al., 2015; O'Connell et al., 2015; Pistilli et al., 2011; Pistilli et al., 2013). Animal experiments have shown that IL-15R $\alpha$  is necessary to maintain insulin sensitivity, since mice lacking IL-15R $\alpha$  are hyperglycemic and insulin-resistant, despite increased oxidative capacity and reduced fat mass (Loro et al., 2015). Furthermore, gene-deletion of IL-15R $\alpha$  in mice is accompanied by enhanced fatigue resistance and a glycolytic-to-oxidative shift in muscle phenotype (O'Connell et al., 2015; Pistilli et al., 2011). It has been demonstrated that strength training promotes a muscle



1 348 myosin heavy chain expression shift from IIx to IIa (Andersen et al., 2005; Campos et al., 2002;  
2  
3 349 Pareja-Blanco et al., 2016), increasing fatigue resistance. However, it remains unknown  
4  
5 350 whether IL-15R $\alpha$  up-regulation contributes to this shift in fiber types (from IIx to IIa) with  
6  
7 351 training in humans. In support of this notion, those participants with a higher IL-15R $\alpha$  protein  
8  
9 352 expression at 4h post-exercise in our study, performed a greater volume of resistance exercise  
10  
11 353 and had a higher baseline leg press 1RM. Thus, the up-regulation of IL-15R $\alpha$  could serve as an  
12  
13 354 adaptive response to maintain muscle characteristics associated with force production. Indeed,  
14  
15 355 others have reported that two SNPs in exon 7 and 4 of the IL-15R $\alpha$  were able to explain a ~11%  
16  
17 356 of the hypertrophy observed after 10 weeks of whole-body resistance exercise in 157 young  
18  
19 357 adults (Riechman et al., 2004). Similarly, another two SNPs, rs2296135 and rs22228059, have  
20  
21 358 been associated with isometric strength and muscle volume before and after 12 weeks of  
22  
23 359 unilateral elbow flexor-extensor resistance exercise (Pistilli et al., 2008).

24  
25 360 Overexpression of IL-15 has revealed that the anabolic/anti-atrophic action of this  
26  
27 361 interleukin is associated with decreased skeletal muscle proteolysis (Busquets et al., 2005) and  
28  
29 362 apoptosis through suppression of DNA fragmentation via tumour necrosis factor alpha (TNF- $\alpha$ )  
30  
31 363 signalling (Figueras et al., 2004). Moreover, IL-15R $\alpha$  mRNA expression is reduced with aging,  
32  
33 364 and may underpin skeletal muscle atrophy in mice (Marzetti et al., 2009). In agreement, we  
34  
35 365 have observed an association between IL-15R $\alpha$  mRNA and MPS in the early recovery phase  
36  
37 366 following resistance exercise. Interestingly, concomitant with the elevation of MPS and IL-  
38  
39 367 15R $\alpha$  mRNA expression, an up-regulation of IL-15R $\alpha$  protein was also found at 4h post-  
40  
41 368 exercise, suggesting that IL-15R $\alpha$  may have a role in the induction or maintenance of the  
42  
43 369 anabolic stimulus during the early post-exercise recovery phase, potentially counteracting the  
44  
45 370 degree of protein breakdown (Phillips et al., 1997). However, further studies are required to  
46  
47 371 delineate the role of IL-15R $\alpha$  in exercise-induced muscle remodelling.

48  
49 372 In contrast to the response observed in skeletal muscle, serum IL-15R $\alpha$  was slightly  
50  
51 373 reduced at 60 min post-exercise. The discordance between the circulating and skeletal muscle  
52  
53 374 IL-15R $\alpha$  response to resistance exercise could imply a counteracting mechanism, by which IL-

15R $\alpha$  binding of IL-15, in blood or cell membrane, reduces its availability (Rubinstein et al., 2006; Schluns et al., 2005) and potentially allows its reabsorption and subsequent restoration of the intracellular pool of free IL-15.

Although pioneer cell culture studies reported an anabolic effect of IL-15 (Quinn et al., 2002; Quinn et al., 1997; Quinn et al., 1995), this has not been confirmed in humans. Strength training-induced muscle hypertrophy is limited to the trained muscles, implying that the anabolic action of IL-15 in human skeletal muscle cannot be explained by an increase in the circulating fraction of this myokine. In fact, human experiments do not give support to an anabolic action of IL-15 in skeletal muscle (Nielsen et al., 2007; Riechman et al., 2004). However, the physiological relevance of the 24h post-exercise elevation of IL-15 gene expression in response to resistance exercise, as reported in the present and previous studies (Nielsen et al., 2007), although suggestive of a role of IL-15 in exercise-induced skeletal muscle adaptation, remains largely unexplained. The lack of association found between the increase in IL-15 mRNA at 24h post-exercise and MPS does not support a critical anabolic role, but to definitely rule out such effect would require the utilization of IL-15 blockers or antibodies, which cannot be used in humans due to potentially intolerable immunological side effects. Moreover, the anti-atrophic effect of IL-15 in skeletal muscle has not been tested in the present study and cannot be excluded (Busquets et al., 2005; Marzetti et al., 2009).

Interleukin-15 is currently considered as a myokine (Grabstein et al., 1994; Quinn et al., 1995). In agreement, we have observed a positive association between serum concentration of IL-15 and IL-15 protein levels in skeletal muscle, suggesting that muscle may be an important source of IL-15 in the basal state. In the present study, we observed that basal IL-15 protein levels in skeletal muscle were associated with serum concentration immediately post-exercise, also suggesting that the size of the intramuscular pool could determine the magnitude of the increase in serum IL-15 elicited by resistance exercise. Nevertheless, the physiological

relevance of the elevated blood IL-15 concentration in close proximity to the end of exercise remains to be elucidated (Riechman et al., 2004; Tamura et al., 2011).

Interestingly, we found that a lower time-under-tension and a lower amount of total weight lifted during resistance exercise session were associated with higher post-exercise serum IL-15 concentration, which could indicate that a more prolonged muscle activation may attenuate the release of IL-15. Depletion of the intracellular pool or reduced *de novo* synthesis of IL-15 could explain the attenuated release of IL-15 with greater training load, given the short life of IL-15 in plasma (Rubinstein et al., 2006; Stoklasek et al., 2006).

The fact that the increase of circulating IL-15 was not accompanied by an increase in its soluble receptor implies that after resistance training there is more free IL-15, available to act on target tissues (Mortier et al., 2004). IL-15 is a potent pro-inflammatory cytokine that stimulates proliferation, maturation and has protective effects on several immune cells (Budagian et al., 2006). In addition to the immunological effects, IL-15 has anti-adipogenic effects in rodents (Carbo et al., 2001). Therefore, although the physiological role of the systemic elevation of IL-15 in response to strength training remains unknown, immunological and metabolic effects are possible.

In conclusion, the IL-15/IL-15R $\alpha$  signalling pathway is activated in human skeletal muscle in response to a single session of resistance exercise. Skeletal muscle mRNA levels and protein IL-15R $\alpha$  expression were elevated four hours after resistance exercise and were positively associated with increased rates of myofibrillar protein synthesis. Therefore, as previously shown in cell culture and *in vivo*, the present investigation lends support to a potentially anabolic effect of IL-15R $\alpha$  in human skeletal muscle. Moreover, our experimental results indicate that IL-15 and IL-15R $\alpha$  may play a role in exercise-induced muscle remodelling. Prolonged resistance training studies are necessary to determine the relevance of IL-15R $\alpha$  in muscle protein synthesis/breakdown, as well as the precise role of circulating and muscular levels of IL-15 and its receptor IL-15R $\alpha$  in chronic physiological adaptations.

428

429 **Perspectives**

430 Previous studies have suggested a role of IL-15R $\alpha$  in muscle phenotypic adaptation to  
431 resistance training. The present study confirms the activation of IL-15/IL-15R $\alpha$  signalling  
432 pathway in human skeletal muscle in response to a single session of resistance exercise. It  
433 remains to be determined how skeletal muscle contributes to circulating levels of IL-15 and  
434 how circulating IL-15 could influence skeletal muscle and adipose tissue mass. Given the  
435 important role that IL-15 has in immune responses, the link between physical activity, skeletal  
436 muscle IL-15 production and immunity deserves further attention. The fact that IL-15R $\alpha$  is  
437 independently associated with myofibrillar protein synthesis and muscle phenotype implies that  
438 the axis IL-15/ IL-15R $\alpha$  may have an important role in human skeletal muscle remodelling.

**ACKNOWLEDGMENTS**

Institutional funding support was provided by the University of Birmingham. APL was recipient of a pre-doctoral fellowship (FPI) from University of Alcalá. JM is supported by the ‘Exercise as Medicine’ PhD scholarship scheme from University of Birmingham. Additional financial support was provided by a Grant from the Ministerio de Economía y Competitividad of Spain (PI14/01509).

The authors would like to thank Alberto Cifuentes for his assistance during qRT-PCR analysis and to Dan Luo, Ismael Pérez-Suarez and Alfredo Santana for their assistance during Western Blot analysis.

**CONFLICT OF INTERESTS**

All the authors declare that they have no conflict of interest derived from the outcomes of this study.

**AUTHOR CONTRIBUTIONS**

APL, JM and LB conceived and designed the experiment. APL, JM and LB collected the data. APL, JM, MMR, DMA, BPK, DV, JB, JALC and LB analysed and interpreted the data. APL, JALC and LB drafted the manuscript and prepared all figures. All authors read and approved the final version of the manuscript.

**REFERENCES**

Andersen LL, Andersen JL, Magnusson SP, Suetta C, Madsen JL, Christensen LR, Aagaard P. Changes in the human muscle force-velocity relationship in response to resistance training and subsequent detraining. *J Appl Physiol* (1985) 2005; 99: 87-94.

Bergstrom J. Percutaneous needle biopsy of skeletal muscle in physiological and clinical research. *Scand J Clin Lab Invest* 1975; 35: 609-616.

- 1 465 Budagian V, Bulanova E, Paus R, Bulfone-Paus S. IL-15/IL-15 receptor biology: a guided tour  
2  
3 466 through an expanding universe. *Cytokine Growth Factor Rev* 2006; 17: 259-280.  
4  
5 467 Busquets S, Figueras MT, Meijssing S, Carbo N, Quinn LS, Almendro V, Argiles JM, Lopez-  
6  
7 468 Soriano FJ. Interleukin-15 decreases proteolysis in skeletal muscle: a direct effect. *Int J Mol*  
8  
9 469 *Med* 2005; 16: 471-476.  
10  
11 470 Campos GE, Luecke TJ, Wendeln HK, Toma K, Hagerman FC, Murray TF, Ragg KE,  
12  
13 471 Ratamess NA, Kraemer WJ, Staron RS. Muscular adaptations in response to three different  
14  
15 472 resistance-training regimens: specificity of repetition maximum training zones. *Eur J Appl*  
16  
17 473 *Physiol* 2002; 88: 50-60.  
18  
19 474 Carbo N, Lopez-Soriano J, Costelli P, Alvarez B, Busquets S, Baccino FM, Quinn LS, Lopez-  
20  
21 475 Soriano FJ, Argiles JM. Interleukin-15 mediates reciprocal regulation of adipose and muscle  
22  
23 476 mass: a potential role in body weight control. *Biochim Biophys Acta* 2001; 1526: 17-24.  
24  
25 477 Dubois S, Mariner J, Waldmann TA, Tagaya Y. IL-15Ralpha recycles and presents IL-15 In  
26  
27 478 trans to neighboring cells. *Immunity* 2002; 17: 537-547.  
28  
29 479 Duitman EH, Orinska Z, Bulanova E, Paus R, Bulfone-Paus S. How a cytokine is chaperoned  
30  
31 480 through the secretory pathway by complexing with its own receptor: lessons from interleukin-  
32  
33 481 15 (IL-15)/IL-15 receptor alpha. *Mol Cell Biol* 2008; 28: 4851-4861.  
34  
35 482 Figueras M, Busquets S, Carbo N, Barreiro E, Almendro V, Argiles JM, Lopez-Soriano FJ.  
36  
37 483 Interleukin-15 is able to suppress the increased DNA fragmentation associated with muscle  
38  
39 484 wasting in tumour-bearing rats. *FEBS Lett* 2004; 569: 201-206.  
40  
41 485 Furmanczyk PS, Quinn LS. Interleukin-15 increases myosin accretion in human skeletal  
42  
43 486 myogenic cultures. *Cell Biol Int* 2003; 27: 845-851.  
44  
45 487 Grabstein KH, Eisenman J, Shanebeck K, Rauch C, Srinivasan S, Fung V, Beers C, Richardson  
46  
47 488 J, Schoenborn MA, Ahdieh M, et al. Cloning of a T cell growth factor that interacts with the  
48  
49 489 beta chain of the interleukin-2 receptor. *Science* 1994; 264: 965-968.  
50  
51 490 Loro E, Seifert EL, Moffat C, Romero F, Mishra MK, Sun Z, Krajacic P, Anokye-Danso F,  
52  
53 491 Summer RS, Ahima RS, Khurana TS. IL-15Ralpha is a determinant of muscle fuel utilization,  
54  
55  
56  
57  
58  
59  
60

1 492 and its loss protects against obesity. *Am J Physiol Regul Integr Comp Physiol* 2015; 309: R835-  
2  
3 493 844.  
4  
5 494 Marzetti E, Carter CS, Wohlgemuth SE, Lees HA, Giovannini S, Anderson B, Quinn LS,  
6  
7 495 Leeuwenburgh C. Changes in IL-15 expression and death-receptor apoptotic signaling in rat  
8  
9 496 gastrocnemius muscle with aging and life-long calorie restriction. *Mech Ageing Dev* 2009; 130:  
10  
11 497 272-280.  
12  
13 498 McKendry J, Perez-Lopez A, McLeod M, Luo D, Dent R, Smeuninx B, Yu J, Taylor AE, Philp  
14  
15 499 A, Breen L. Short inter-set rest blunts resistance exercise-induced increases in myofibrillar  
16  
17 500 protein synthesis and intracellular signaling in young males. *Exp Physiol* 2016; 101: 866-882.  
18  
19 501 Mortier E, Bernard J, Plet A, Jacques Y. Natural, proteolytic release of a soluble form of human  
20  
21 502 IL-15 receptor alpha-chain that behaves as a specific, high affinity IL-15 antagonist. *J Immunol*  
22  
23 503 2004; 173: 1681-1688.  
24  
25 504 Nielsen AR, Mounier R, Plomgaard P, Mortensen OH, Penkowa M, Speerschneider T,  
26  
27 505 Pilegaard H, Pedersen BK. Expression of interleukin-15 in human skeletal muscle effect of  
28  
29 506 exercise and muscle fibre type composition. *J Physiol* 2007; 584: 305-312.  
30  
31 507 O'Connell G, Guo G, Stricker J, Quinn LS, Ma A, Pistilli EE. Muscle-specific deletion of exons  
32  
33 508 2 and 3 of the IL15RA gene in mice: effects on contractile properties of fast and slow muscles.  
34  
35 509 *Journal of applied physiology* 2015; 118: 437-448.  
36  
37 510 O'Connell GC, Nichols C, Guo G, Croston TL, Thapa D, Hollander JM, Pistilli EE. IL-  
38  
39 511 15Ralpha deficiency in skeletal muscle alters respiratory function and the proteome of  
40  
41 512 mitochondrial subpopulations independent of changes to the mitochondrial genome.  
42  
43 513 *Mitochondrion* 2015; 25: 87-97.  
44  
45 514 Pareja-Blanco F, Rodriguez-Rosell D, Sanchez-Medina L, Sanchis-Moysi J, Dorado C, Mora-  
46  
47 515 Custodio R, Yanez-Garcia JM, Morales-Alamo D, Perez-Suarez I, Calbet JA, Gonzalez-Badillo  
48  
49 516 JJ. Effects of velocity loss during resistance training on athletic performance, strength gains and  
50  
51 517 muscle adaptations. *Scand J Med Sci Sports* 2016.



1 518 Phillips SM, Tipton KD, Aarsland A, Wolf SE, Wolfe RR. Mixed muscle protein synthesis and  
2  
3 519 breakdown after resistance exercise in humans. *Am J Physiol* 1997; 273: E99-107.  
4  
5 520 Philp A, Chen A, Lan D, Meyer GA, Murphy AN, Knapp AE, Olfert IM, McCurdy CE,  
6  
7 521 Marcotte GR, Hogan MC, Baar K, Schenk S. Sirtuin 1 (SIRT1) deacetylase activity is not  
8  
9 522 required for mitochondrial biogenesis or peroxisome proliferator-activated receptor-gamma  
10  
11 523 coactivator-1alpha (PGC-1alpha) deacetylation following endurance exercise. *J Biol Chem*  
12  
13 524 2011; 286: 30561-30570.  
14  
15 525 Pistilli EE, Bogdanovich S, Garton F, Yang N, Gulbin JP, Conner JD, Anderson BG, Quinn LS,  
16  
17 526 North K, Ahima RS, Khurana TS. Loss of IL-15 receptor alpha alters the endurance,  
18  
19 527 fatigability, and metabolic characteristics of mouse fast skeletal muscles. *J Clin Invest* 2011:  
20  
21 528 121: 3120-3132.  
22  
23 529 Pistilli EE, Devaney JM, Gordish-Dressman H, Bradbury MK, Seip RL, Thompson PD,  
24  
25 530 Angelopoulos TJ, Clarkson PM, Moyna NM, Pescatello LS, Visich PS, Zoeller RF, Gordon  
26  
27 531 PM, Hoffman EP. Interleukin-15 and interleukin-15R alpha SNPs and associations with muscle,  
28  
29 532 bone, and predictors of the metabolic syndrome. *Cytokine* 2008; 43: 45-53.  
30  
31 533 Pistilli EE, Guo G, Stauber WT. IL-15Ralpha deficiency leads to mitochondrial and myofiber  
32  
33 534 differences in fast mouse muscles. *Cytokine* 2013; 61: 41-45.  
34  
35 535 Pistilli EE, Siu PM, Alway SE. Interleukin-15 responses to aging and unloading-induced  
36  
37 536 skeletal muscle atrophy. *Am J Physiol Cell Physiol* 2007; 292: C1298-1304.  
38  
39 537 Quinn LS, Anderson BG, Drivdahl RH, Alvarez B, Argiles JM. Overexpression of interleukin-  
40  
41 538 15 induces skeletal muscle hypertrophy in vitro: implications for treatment of muscle wasting  
42  
43 539 disorders. *Exp Cell Res* 2002; 280: 55-63.  
44  
45 540 Quinn LS, Haugk KL, Damon SE. Interleukin-15 stimulates C2 skeletal myoblast  
46  
47 541 differentiation. *Biochem Biophys Res Commun* 1997; 239: 6-10.  
48  
49 542 Quinn LS, Haugk KL, Grabstein KH. Interleukin-15: a novel anabolic cytokine for skeletal  
50  
51 543 muscle. *Endocrinology* 1995; 136: 3669-3672.  
52  
53  
54  
55  
56  
57  
58  
59  
60



1 544 Riechman SE, Balasekaran G, Roth SM, Ferrell RE. Association of interleukin-15 protein and  
2  
3 545 interleukin-15 receptor genetic variation with resistance exercise training responses. J Appl  
4  
5 546 Physiol (1985) 2004: 97: 2214-2219.  
6  
7 547 Rubinstein MP, Kovar M, Purton JF, Cho JH, Boyman O, Surh CD, Sprent J. Converting IL-15  
8  
9 548 to a superagonist by binding to soluble IL-15R{alpha}. Proceedings of the National Academy  
10  
11 549 of Sciences of the United States of America 2006: 103: 9166-9171.  
12  
13 550 Sato N, Patel HJ, Waldmann TA, Tagaya Y. The IL-15/IL-15Ralpha on cell surfaces enables  
14  
15 551 sustained IL-15 activity and contributes to the long survival of CD8 memory T cells. Proc Natl  
16  
17 552 Acad Sci U S A 2007: 104: 588-593.  
18  
19 553 Schluns KS, Stoklasek T, Lefrancois L. The roles of interleukin-15 receptor alpha: trans-  
20  
21 554 presentation, receptor component, or both? The international journal of biochemistry & cell  
22  
23 555 biology 2005: 37: 1567-1571.  
24  
25 556 Stoklasek TA, Schluns KS, Lefrancois L. Combined IL-15/IL-15Ralpha immunotherapy  
26  
27 557 maximizes IL-15 activity in vivo. J Immunol 2006: 177: 6072-6080.  
28  
29 558 Tamura Y, Watanabe K, Kantani T, Hayashi J, Ishida N, Kaneki M. Upregulation of circulating  
30  
31 559 IL-15 by treadmill running in healthy individuals: is IL-15 an endocrine mediator of the  
32  
33 560 beneficial effects of endurance exercise? Endocr J 2011: 58: 211-215.  
34  
35 561 Wolfe RR, Chinkes DL. *Isotope Tracers in Metabolic Research: Principles and Practice of*  
36  
37 562 *Kinetic Analysis*: Wiley 2005.  
38  
39  
40  
41  
42 563  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

## FIGURE LEGENDS

**Figure 1.** IL-15 response to a single session of resistance exercise.

IL-15 mRNA (qRT-PCR) (A) and protein levels (Western blotting) (B) from *vastus lateralis* muscle biopsies, and serum IL-15 levels (ELISA) (C). Values are presented as means  $\pm$  SD (N = 14). \* P<0.05 compared to Pre-exercise. # P<0.05 compared to 0h post-exercise. Data were log-transformed.

**Figure 2.** IL-15R $\alpha$  response to a single session of resistance exercise.

IL-15R $\alpha$  mRNA (qRT-PCR) (A) and protein levels (Western blotting) (B) from *vastus lateralis* muscle biopsies, and serum IL-15R $\alpha$  levels (ELISA) (C). Values are presented as means  $\pm$  SD (N = 14). \* P<0.05 compared to Pre-exercise. # P<0.05 compared to 0h post-exercise.  $\Phi$  P<0.05 compared to 24h post-exercise. § P<0.05 differences compared to Mid-exercise. Data were log-transformed.

**Figure 3.** Relationship between myofibrillar protein synthesis measured as a fractional synthetic rate (FSR) and IL-15R $\alpha$  mRNA pre- and post-exercise. The association remained significant in Fig. 3B ( $r = 0.665$ ,  $P = 0.026$ ) when the lowest FSR values were excluded. a.u., arbitrary units.

**Supplementary Figure 1.** Skeletal muscle IL-15 response to a single session of resistance exercise by group (1 and 5 min groups).

IL-15 mRNA (qRT-PCR) (A) and protein levels (Western blotting) (B) from *vastus lateralis* muscle biopsies. Values are presented as means  $\pm$  SD (N = 14). \* P<0.05 compared to Pre-exercise. Data were log-transformed. a.u., arbitrary units.

1  
2 589 **Supplementary Figure 2.** Skeletal muscle IL-15R $\alpha$  response to a single session of resistance  
3  
4 590 exercise by group (1 and 5 min groups).  
5  
6 591 IL-15R $\alpha$  mRNA (qRT-PCR) (A) and protein levels (Western blotting) (B) from *vastus lateralis*  
7  
8 592 muscle biopsies. Values are presented as means  $\pm$  SD (N = 14). \* P<0.05 compared to Pre-  
9  
10 593 exercise. # P<0.05 compared to 0h post-exercise. Data were log-transformed. a.u., arbitrary  
11  
12  
13 594 units.  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

**Table 1.** Participants' physical characteristics

N	14
Age (yrs)	24.9 ± 4.8
Body mass (kg)	82.2 ± 11.9
BMI (kg/m <sup>2</sup> )	25.6 ± 3.1
Whole-body FFM (kg)	66.0 ± 8.8
Legs FFM (kg)	21.5 ± 3.2
Whole-Body FM (kg)	12.9 ± 5.2
Legs FM (kg)	4.5 ± 1.5
Leg Press 1-RM (kg)	268 ± 51
Knee Extension (kg)	169 ± 26
Training experience (yrs)	6 ± 5
Leg training (days/week)	2 ± 1

Values are presented as mean ± SD. BMI, body mass index; FFM, fat free mass; FM, fat mass; 1RM, one-repetition maximum.

**Table 2.** Primers for qRT-PCR analysis.

Primer	Sequence	Accession Number	T <sub>m</sub>
IL-15	F: 5'-AAAGTGATGTTCAACCCAGTTG	NM_000585.4	60°
	R: 3'-CCTCCAGTTCCTCACATTCTTTG		30s
IL-15Rα	F: 5'-CAGCCGCCAGGTGTGTATC	NM_002189.3	60°
	R: 3'-TTGCCTTGACTTGAGGTAGCA		30s

T<sub>m</sub>, melting temperature.

1 **Table 3.** Variables describing the resistance session.

	Leg Press					Knee Extension					Total
Set	1	2	3	4	Total	1	2	3	4	Total	8
Load (kg)	205±40	205±39	205±38	206±37	820±154	123±18	119±19	117±19	114±18	473±73	1293±210
Repetition	13±3	11±2	11±2	10±2	44±6	9±2	9±2	10±2	9±2	37±6	10±1
Volume (kg)	2484±401	2188±296	2260±452	2019±533	8951±1217	1068±276	1119±296	1139±323	1081±352	4407±1127	13358±2026
T-U-T (s)	38.3±11.5	34.4±7.1	33.1±6.2	32.9±5.9	138.7±27.4	23.7±9.5	18.8±3.4	19.2±4.7	18.9±3.1	80.6±17.2	219.4±40.8
RPE (0-10)	9±1	9±1	10±0	10±1	9±0	10±0	10±0	10±0	10±0	10±0	10±0

2 Values are presented as mean ± SD. T-U-T, time-under-tension; RPE, rating of perceived exertion.











